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		To:	Examiner Kinsey White, N.E TC/AU: 1648					
	I	Firm:	USPTO					
ļ	Facsimile	No.:	(571) 273-9943					
		rom:	Mary J. Wilson					
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,	ATTACHMENT	/S: D	RAFT SET O	F PROPOS	SED REVI	SED CLAIMS		
	in re Patent Ap	re Patent Application of:						
	HELLSTROM e Serial No. 10/5 Filed: May 10, For: METHO	78,848 2006	D MEANS RE	LATING T	O HEPATI	TIS B INFEÇT	ΓΙΟΝ	
	Dear Examiner		N.					
\ 	vith reference to our telephone discussion of earlier, attached is a copy of the proposed vised claims for your review.							
İ	look forward to speaking with you on Thursday.							
E	Best regards.	:						
Ŋ	Mary J. Wilson	:						

CONFIDENTIALITY NOTE

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- 1. (Currently Amended) A method of predicting determining whether an individual having hepatitis B virus (HBV) infection will respond to interferon alpha (IFNα) treatment, the method comprising:
 - i) obtaining a pre-treatment sample from said HBV-infected individual.
- analyzing said pre-treatment sample for determining the presence or absence of antibodies reactive with a preS1 peptide consisting of the sequence of residues 94-117 (SEQ ID NO:1) in a pre-treatment sample obtained from the individual, and;

predicting from wherein the presence of said antibodies in said <u>pre-treatment</u> sample <u>indicates</u> that said individual will respond to said treatment or <u>predicting from and</u> the absence of said antibodies in said <u>pre-treatment</u> sample <u>indicates</u> that said individual will not respond to said treatment.

- 2.-3. (Cancelled).
- 4. (Previously Presented) A method according to claim 1 wherein the individual has chronic HBV infection.
- 5. (Previously Presented) A method according to claim 1 wherein the individual is HBeAg positive.
- 6. (Previously Presented) A method according to claim 1 wherein the individual is HBeAg negative.

- 7. (Currently Amended) A method according to claim 1 wherein, in step (ii), said pre-treatment sample is analyzed for the presence or absence of the antibodies are IgG or IgM antibodies reactive with said preS1 peptide.
- 8. (Previously Presented) A method according to claim 1 wherein the pre-treatment sample is a blood, serum or plasma sample.
- 9. (Currently Amended) A method according to claim 1 wherein step (ii) comprises comprising;

contacting the pre-treatment sample with a preS1 peptide consisting of the sequence of residues 94-117 (SEQ ID NO:1) and;

detecting the presence or absence of binding of determining binding of said antibodies to said preS1 peptide.

- 10. (Currently Amended) A method according to claim 9 wherein the <u>preS1</u> peptide comprises a detectable label.
- 11. (Currently Amended) A method according to claim 9 wherein said <u>preS1</u> peptide is immobilised.
- 12. (Previously Presented) A method according to claim 9 wherein said binding is detected with a labelled secondary antibody.

- 20. (Currently Amended) A method of treating a hepatitis B an HBV infection in an individual identified as being responsive to IFNa comprising:
 - i) obtaining a pre-treatment sample from an HBV-infected individual,
- <u>ii)</u> analyzing said pre-treatment sample for determining the presence or absence of antibodies reactive with a preS1 peptide consisting of the sequence of residues 94-117 (SEQ ID NO:1) in a pre-treatment sample obtained from the individual;

predicting from wherein the presence of said antibodies in said pre-treatment sample indicates that said individual will respond to interferon alpha (IFN α) IFN α treatment, and;

administering IFN α to said individual identified as being responsive to IFN α .

- 21. (Cancelled).
- 22. (Currently Amended) A method according to claim 20 <u>further comprising</u> administering wherein corticosteroid to said individual identified as being responsive to IFNa is administered to the individual.
- 23. (Cancelled).